

Rec'd PCT/PTO 01 DEC 2004
Ref 3002450

GB 03/2438

INVESTOR IN PEOPLE



PRIORITY DOCUMENT

SUBMITTED OR TRANSMITTED IN
COMPLIANCE WITH RULE 17.1(a) OR (b)

The Patent Office
Concept House
Cardiff Road

Newport
South Wales
NP10 8QQ

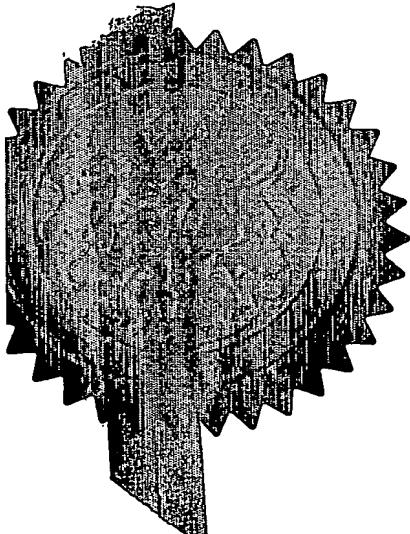
REC'D 03 JUL 2003
WPO PCT

I, the undersigned, being an officer duly authorised in accordance with Section 74(1) and (4) of the Deregulation & Contracting Out Act 1994, to sign and issue certificates on behalf of the Comptroller-General, hereby certify that annexed hereto is a true copy of the documents as originally filed in connection with the patent application identified therein.

In accordance with the Patents (Companies Re-registration) Rules 1982, if a company named in this certificate and any accompanying documents has re-registered under the Companies Act 1980 with the same name as that with which it was registered immediately before re-registration save for the substitution as, or inclusion as, the last part of the name of the words "public limited company" or their equivalents in Welsh, references to the name of the company in this certificate and any accompanying documents shall be treated as references to the name with which it is so re-registered.

In accordance with the rules, the words "public limited company" may be replaced by p.l.c., plc, P.L.C. or PLC.

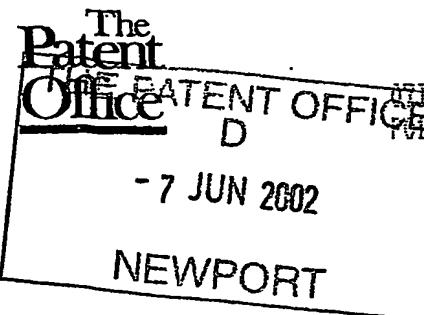
Re-registration under the Companies Act does not constitute a new legal entity but merely subjects the company to certain additional company law rules.



Signed *R. Mahoney*
Dated 25 June 2003

Request for grant of a patent

(See the notes on the back of this form. You can also get an explanatory leaflet from the Patent Office to help you fill in this form)



The Patent Office

Cardiff Road
Newport
Gwent NP9 1RH

1. Your reference

SMC 60525/GB/PI

2. Patent application number

(The Patent Office will fill in this part)

0213011.0

57 JUN 2002

3. Full name, address and postcode of the or of each applicant (underline all surnames)

Avecia Limited
Hexagon House
Blackley
Manchester, M9 8ZS
United Kingdom

Patents ADP number (if you know it)

07764137001

If the applicant is a corporate body, give the country/state of its incorporation

GB

4. Title of the invention

COMPOUNDS

5. Name of your agent (if you have one)

MAYALL, John

"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)

Avecia Limited
Hexagon House
Blackley
Manchester, M9 8ZS
United Kingdom

Patents ADP number (if you know it)

62443130024

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (if you know it) the or each application number

Country Priority application number
(if you know it) Date of filing
(day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application Date of filing
(day / month / year)

8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if:

- a) any applicant named in part 3 is not an inventor, or
- b) there is an inventor who is not named as an applicant, or
- c) any named applicant is a corporate body.

See note (d))

Patents Form 1/77

9. Enter the number of sheets for any of the following items you are filing with this form.
Do not count copies of the same document

Continuation sheets of this form

Description	19
Claim(s)	02
Abstract	
Drawing(s)	

8

10. If you are also filing any of the following, state how many against each item.

Priority documents

Translations of priority documents

Statement of inventorship and right to grant of a patent (Patents Form 7/77)

Request for preliminary examination and search (Patents Form 9/77)

Request for substantive examination
(Patents Form 10/77)

Any other documents
(please specify)

11. I/We request the grant of a patent on the basis of this application.

Signature

G. Terry

Date 6/6/02

Avecia Limited Authorised Signatory

12. Name and daytime telephone number of person to contact in the United Kingdom

Mrs K.M. Pinder/Miss G. Terry 0161 721 1361/2

Warning

After an application for a patent has been filed, the Comptroller of the Patent Office will consider whether publication or communication of the invention should be prohibited or restricted under Section 22 of the Patents Act 1977. You will be informed if it is necessary to prohibit or restrict your invention in this way. Furthermore, if you live in the United Kingdom, Section 23 of the Patents Act 1977 stops you from applying for a patent abroad without first getting written permission from the Patent Office unless an application has been filed at least 6 weeks beforehand in the United Kingdom for a patent for the same invention and either no direction prohibiting publication or communication has been given, or any such direction has been revoked.

Notes

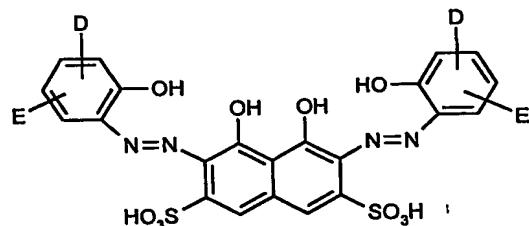
- a) If you need help to fill in this form or you have any questions, please contact the Patent Office on 0645 500505.
- b) Write your answers in capital letters using black ink or you may type them.
- c) If there is not enough space for all the relevant details on any part of this form, please continue on a separate sheet of paper and write "see continuation sheet" in the relevant part(s). Any continuation sheet should be attached to this form.
- d) If you have answered 'Yes' Patents Form 7/77 will need to be filed.
- e) Once you have filled in the form you must remember to sign and date it.
- f) For details of the fee and ways to pay please contact the Patent Office.

COMPOUNDS

This invention relates to compounds suitable for use as dyes and processes for their manufacture, to compositions derived therefrom and to their use in printing, especially ink jet printing ("IJP"). IJP is a non-impact printing technique in which droplets of a composition are ejected through a fine nozzle onto a substrate without bringing the nozzle into contact with the substrate.

There are many demanding performance requirements for dyes and compositions used in IJP. For example they desirably provide sharp, non-feathered images having good water-fastness, light-fastness and optical density. The compositions are often required to dry quickly when applied to a substrate to prevent smudging, but they should not form a crust over the tip of an ink jet nozzle because this will stop the printer from working. The compositions should also be stable to storage over time without decomposing or forming a precipitate which could block the fine nozzle.

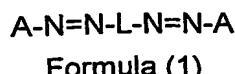
GB 1,130,348 discloses the preparation of metallised disazo reactive dyes derived from chromotropic acid for the conventional dyeing of textiles. US 4,361,842 discloses the use of metallised compounds derived from chromotropic acid to prolong the service life of recording heads in thermal ink jet printers. Japanese application JP 2002-20659 discloses recording liquids for ink-jet printing comprising metallised dyes containing a group of the following formula:



wherein D and E are as defined in JP 2002-20659. In the above dyes of JP2002-20659 both of the phenylazo groups attached to the central chromotropic acid group have a hydroxy substituent ortho to the -N=N- groups shown above.

Surprisingly it has been found that dyes lacking at least one of the ortho hydroxy groups in the above formula have outstanding properties as ink jet inks. Furthermore, dyes lacking metallisation of the 1,8-dihydroxy groups in the central chromotropic acid group are also valuable colorants for ink jet printing.

According to the present invention there is provided a process for printing an image on a substrate comprising applying thereto a composition comprising a liquid medium and a compound of Formula (1):



wherein:

each A independently is optionally substituted aryl or heteroaryl; and

L is an optionally substituted, optionally metallised 1,8-dihydroxynaphthylene group;

provided that at most one of the groups represented by A has a hydroxy substituent ortho to the -N=N- groups shown in Formula (1).

Preferably the composition is applied to the substrate by means of an ink jet printer. The ink jet printer preferably applies the composition to the substrate in the form of droplets which are ejected through a small orifice onto the substrate. Preferred ink jet printers are piezoelectric ink jet printers and thermal ink jet printers. In thermal ink jet printers, programmed pulses of heat are applied to the composition in a reservoir by means of a resistor adjacent to the orifice, thereby causing the composition to be ejected in the form of small droplets directed towards the substrate during relative movement between the substrate and the orifice. In piezoelectric ink jet printers the oscillation of a small crystal causes ejection of the composition from the orifice.

The substrate is preferably paper, plastic, a textile, metal or glass, more preferably a treated substrate such as a coated paper or coated plastic, especially coated paper.

Preferred plain or treated papers are papers which may have an acid, alkaline or neutral character. Examples of commercially available treated papers include HP PremiumTM Coated Paper, HP PhotopaperTM (both available from Hewlett Packard Inc.); Stylus ProTM 720 dpi Coated Paper, Epson Photo QualityTM Glossy Film, Epson Photo QualityTM Glossy Paper (all available from Seiko Epson Corp.); Canon HR 101TM High Resolution Paper, Canon GP 201TM Glossy Paper, Canon HG 101TM and HG 201TM High Gloss Film, Canon PR101TM (all available from Canon); Kodak Premium PhotopaperTM, Kodak Premium InkJetpaperTM (available from Kodak); Konica Inkjet Paper QPTM Professional Photo Glossy, Konica Inkjet Paper QPTM Professional Photo 2-sided Glossy, Konica Inkjet Paper QPTM Premium Photo Glossy, Konica Inkjet Paper QPTM Premium Photo Silky (available from Konica).

Preferred optionally substituted aryl groups represented by A are optionally substituted phenyl and naphthyl. Preferred optionally substituted heteroaryl groups represented by A are any heterocycle or substituted heterocycle comprising a 5 to 7 membered ring, more preferably optionally substituted pyridyl, pyrazolyl or 1,2,4-triazole.

The groups represented by A in Formula (1) may be the same as each other or, more preferably, different from each other.

Optional substituents which may be present on A are preferably selected from OH (subject to the proviso that at most one of the groups represented by A has a hydroxy substituent ortho to the -N=N- groups shown in Formula (1)), SO₃H, PO₃H₂, CO₂H, NO₂, NH₂, optionally substituted alkyl (especially C₁₋₄-alkyl optionally carrying a sulpho, carboxy, phosphato, C₁₋₄-alkoxy, amino or hydroxy group), optionally substituted alkoxy (especially C₁₋₄-alkoxy optionally carrying a sulpho, carboxy, phosphato, C₁₋₄-alkoxy, C₁₋₄-alkyl, amino or hydroxy group), amine (especially N carrying one or two C₁₋₄-alkyl groups optionally carrying a sulpho, carboxy, phosphato, C₁₋₄-alkoxy, amino or hydroxy group),

acylamine (especially C₁₋₄-acylamino) and ureido. Preferred substituents for A are sulpho, hydroxy, nitro, amino, acetyl amino, carboxy and 3-carboxy-1,2,4-triazole.

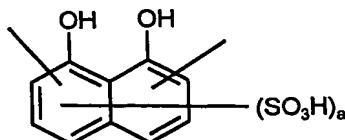
Preferably each A independently carries from 0 to 3 substituents, more preferably 1 or 2 substituents.

As examples of optionally substituted phenyl groups represented by A there may be mentioned 2-sulphophenyl, 3-sulfophenyl, 4-sulfophenyl, 2-nitrophenyl, 3-nitrophenyl, 4-nitrophenyl, 2-sulfo-4-phosphatophenyl, 2-sulfo-4-aminophenyl, 2-sulfo-4-acetylaminophenyl, 2-sulfo-4-methoxyphenyl, 2-sulfo-5-aminophenyl, 3-sulfo-4-nitrophenyl, 3-sulfo-4-aminophenyl, 2-sulfo-4-nitrophenyl, 2,5-disulfophenyl, 2,5-disulfo-4-acetylaminophenyl, 2-hydroxy-3,5-disulfophenyl, 2-carboxy-4-acetylaminophenyl, 2-carboxy-4-aminophenyl, 3,5-dicarboxyphenyl and 2,5-dihydroxyethoxyphenyl.

Preferably both of the groups represented by A are free from hydroxy substituents ortho to the -N=N- groups shown in Formula (1).

L is preferably optionally substituted 1,8-dihydroxynaphthylene, i.e. unmetallised.

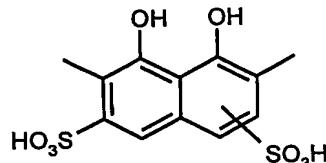
Preferably L is of Formula (2) or a metal complex thereof:



Formula (2)

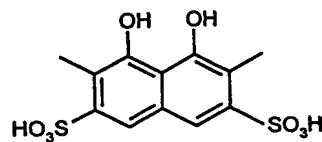
wherein a is 1 or 2 and SO₃H is in free acid or salt form. Preferably a is 2 and the SO₃H groups shown in Formula (2) are in the 3- and 6- positions or the 3- and 5- positions.

In a preferred aspect of the invention, L is of Formula (3) or a metal complex thereof:

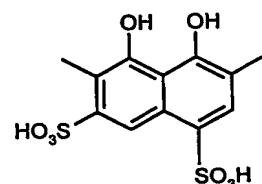


Formula (3)

More preferably L is of Formula (4) or (5) or a metal complex thereof:



Formula (4)



Formula (5)

In Formulae (3), (4) and (5) the SO₃H groups include the free acid and salt forms.

When L is a metallised 1,8-dihydroxynaphthylene group (i.e. a metal complex) the metal is preferably B or a transition metal, more preferably Mn, Fe, Cr, Co, Ni, Cu or Zn, especially Co, Ni or Cu. The metal may be complexed with the compound of Formula (1) in a ratio of from 1:2 to 2:1, preferably in a ratio of metal to compound of Formula (1) of 1:2, 2:3, 1:1, 2:2 or 2:1, especially 2:1. However we have found that when L is not metallised the compound of Formula (1) is still a valuable colorant for ink jet printing.

Such unmetallised dyes are cheaper and easier to make than the corresponding metallised dyes and they are more environmentally friendly due to the absence of

transition metals. Furthermore the absence of transition metals also reduces the likelihood of nozzle blockage in thermal ink jet printers.

According to a second aspect of the present invention there is provided a composition comprising:

(a) 0.1 to 15 parts of the compound of Formula (1) as hereinbefore defined; and

(b) from 85 to 99.9 parts of a liquid medium;

wherein all parts are by weight and the number of parts of (a)+(b)=100.

The composition used in the first aspect of the invention is preferably as defined in the second aspect of the present invention.

The number of parts of component (a) is preferably from 0.2 to 12, more preferably from 0.5 to 10, and especially from 1 to 8 parts. The number of parts of component (b) is preferably from 99.8 to 88, more preferably from 99.5 to 90, especially from 99 to 92 parts.

When the medium is a liquid, preferably component (a) is completely dissolved in component (b). Preferably component (a) has a solubility in component (b) at 20°C of at least 10%. This allows the preparation of liquid dye concentrates which may be used to prepare inks and reduces the chance of the dye precipitating if evaporation of the liquid medium occurs during storage.

Preferred liquid media include water or a mixture of water and an organic solvent.

When the liquid medium comprises a mixture of water and an organic solvent, the weight ratio of water to organic solvent is preferably from 99:1 to 1:99, more preferably from 99:1 to 50:50 and especially from 95:5 to 70:30.

It is preferred that the organic solvent present in the mixture of water and organic solvent is a water-miscible organic solvent or a mixture of such solvents. Preferred water-miscible organic solvents include C₁₋₆-alkanols, preferably methanol, ethanol, n-propanol, isopropanol, n-butanol, sec-butanol, tert-butanol, n-pentanol, cyclopentanol and cyclohexanol; linear amides, preferably dimethylformamide or dimethylacetamide; ketones and ketone-alcohols, preferably acetone, methyl ether ketone, cyclohexanone and diacetone alcohol; water-miscible ethers, preferably tetrahydrofuran and dioxane; diols, preferably diols having from 2 to 12 carbon atoms, for example pentane-1,5-diol, ethylene glycol, propylene glycol, butylene glycol, pentylene glycol, hexylene glycol and thiodiglycol and oligo- and poly-alkyleneglycols, preferably diethylene glycol, triethylene glycol,

polyethylene glycol and polypropylene glycol; triols, preferably glycerol and 1,2,6-hexanetriol; mono-C₁₋₄-alkyl ethers of diols, preferably mono-C₁₋₄-alkyl ethers of diols having 2 to 12 carbon atoms, especially 2-methoxyethanol, 2-(2-methoxyethoxy)ethanol, 2-(2-ethoxyethoxy)-ethanol, 2-[2-(2-methoxyethoxy)ethoxy]ethanol, 2-[2-(2-ethoxyethoxy)-ethoxy]-ethanol and ethyleneglycol monoallylether; cyclic amides, preferably 2-pyrrolidone, N-methyl-2-pyrrolidone, N-ethyl-2-pyrrolidone, caprolactam and 1,3-dimethylimidazolidone; cyclic esters, preferably caprolactone; sulphoxides, preferably dimethyl sulphoxide and sulpholane. Preferably the liquid medium comprises water and 2 or more, especially from 2 to 8, water-soluble organic solvents.

Especially preferred water-soluble organic solvents are cyclic amides, especially 2-pyrrolidone, N-methyl-pyrrolidone and N-ethyl-pyrrolidone; diols, especially 1,5-pentane diol, ethyleneglycol, thiodiglycol, diethyleneglycol and triethyleneglycol; and mono- C₁₋₄-alkyl and C₁₋₄-alkyl ethers of diols, more preferably mono- C₁₋₄-alkyl ethers of diols having 2 to 12 carbon atoms, especially ((2-methoxy-2)-ethoxy)-2-ethoxyethanol.

Examples of further suitable ink media comprising a mixture of water and one or more organic solvents are described in US 4,963,189, US 4,703,113, US 4,626,284 and EP 4,251,50A.

The pH of the composition is preferably from 4 to 11, more preferably from 7 to 10.

The viscosity of the composition at 25°C is preferably less than 50cP, more preferably less than 20 cP and especially less than 5cP.

When the compositions according to the invention are used as ink jet printing compositions, the composition preferably has a concentration of halide ions of less than 500 parts per million, more preferably less than 100 parts per million. It is especially preferred that the composition has less than 100, more preferably less than 50 parts per million of divalent and trivalent metals, wherein parts refer to parts by weight relative to the total weight of the composition. We have found that purifying the compositions to reduce the concentration of these undesirable ions reduces nozzle blockage in ink jet printing heads, particularly in thermal ink jet printers.

The compounds of Formula (1) may be used as the sole colorant in the compositions because of their attractive black shade. However, if desired, one may combine the present compounds with one or more further colorants if a slightly different shade is required for a particular end use. The further colorants are preferably dyes. When further colorants are included in the composition these are preferably selected from but not limited to black, magenta, cyan and yellow colorants and combinations thereof.

Suitable further black colorants include C.I.Food Black 2, C.I.Direct Black 19, C.I.Reactive Black 31, PRO-JET™ Fast Black 2, C.I.Direct Black 195; C.I.Direct Black 168; and black dyes described in patents by Lexmark (e.g. EP 0 539,178 A2, Example 1, 2, 3, 4 and 5), Orient Chemicals (e.g. EP 0 347 803 A2, pages 5-6, azo dyes 3, 4, 5, 6, 7, 8, 12, 13, 14, 15 and 16) and Seiko Epson Corporation.

Suitable further magenta colorants include PRO-JET™ Fast Magenta 2.

Suitable further yellow colorants include C.I.Direct Yellow 142; C.I.Direct Yellow 132; C.I.Direct Yellow 86; C.I.Direct Yellow 85; C.I. Direct Yellow 173; and C.I.Acid Yellow 23.

Suitable further cyan colorants include phthalocyanine colorants, Direct Blue 199 and Acid Blue 99.

The composition according to the second aspect of the present invention and that used in the process according to the first aspect of the present invention may also contain additional components conventionally used in ink jet printing inks, for example viscosity and surface tension modifiers, corrosion inhibitors, biocides, kogation reducing additives and surfactants which may be ionic or non-ionic.

In a third aspect of the invention we have also devised a process for preparing a compound of Formula (1) as hereinbefore defined which comprises diazotising an amine of formula A-NH₂ to give a diazonium salt, and coupling the resultant diazonium salt with a compound of Formula (6):



Formula (6)

wherein L and each A independently is as hereinbefore defined.

Preferably the diazotisation is performed using a diazotising agent, especially sodium nitrite. Further preferably the diazotisation is performed at a temperature of 0 to 5°C. Surprisingly we have found that the coupling can be performed under conditions of moderate, i.e relatively neutral, pH facilitated by the use of substantial amounts of aprotic solvents. Typically the diazotisation is performed in a liquid medium. Preferred liquid media are water and compositions comprising water and an organic solvent, especially an aprotic solvent. Aprotic solvents which may be used may be chosen from, for example, dimethylformamide, N-methylpyrrolidone, 2-pyrrolidone, ethers, acetone, glymes such as polyglyme (MW 300) or polyethylene glycols and/or mixtures thereof. Particularly preferred aprotic solvents are tetrahydrofuran and polyglyme. The aprotic solvent is preferably present in the range 0 to 80 wt%, preferably 20 to 60 wt%, especially 25 to 45 wt% relative to the total weight of the liquid medium. Use of substantial amounts of aprotic solvents has the advantage of permitting diazo coupling at much lower pH than customary (usually pH in the region of 14), thereby significantly improving the efficiency, from 30% to as much as 70%, of the process as a result of greatly reduced decomposition of diazonium salts.

Preferably the process of the third aspect of the present invention is performed in the presence of an acid binding agent to maintain a moderate pH. Preferably the acid binding agent used is any which maintains the pH between 5 and 9, preferably between 6 and 7.5. Further, the acid binding agent is preferably selected from but is not limited to alkali metal hydroxides, carbonates, bicarbonates or phosphates or organic bases such as triethanolamine or triethylamine. Particularly preferred alkali metal hydroxide acid-binding agents are NaOH, KOH and LiOH, especially NaOH and LiOH. Particularly preferred alkali

metal carbonate acid-binding agents are Li_2CO_3 and Na_2CO_3 , especially Na_2CO_3 . This has the advantage of maintaining pH sufficiently low to reduce the decomposition of diazonium salts significantly.

5 A fourth feature of the present invention comprises a process for the preparation of a compound of Formula-(1) which comprises the reaction of a compound of formula A-
N=N-Q-N=N-A (wherein each A independently is as hereinbefore defined and Q is an
optionally substituted 1-hydroxy-8-amino-naphthylene group) with a strong base thereby
forming a compound for Formula (1) as hereinbefore defined, optionally followed by
10 contacting the compound so formed with a metal salt. Preferably the strong base is a
metal hydroxide, more preferably an alkali metal hydroxide, especially sodium or
potassium hydroxide. The reaction with strong base preferably comprises heating in
solution at an elevated temperature until reaction is complete. Preferred elevated
temperatures are 60 to 90 °C, more preferably 65 to 80 °C, especially 65 to 75 °C. The pH
is preferably alkaline, more preferably 10 to 14, especially 12 to 14.

15 The compound of formula A-N=N-Q-N=N-A is preferably formed by diazotising an
amine of formula A-NH₂, as defined above, and coupling the resultant diazonium salt with
an azo compound of Formula H-Q-N=N-A wherein each A independently and Q are
hereinbefore defined.

20 The process according to the fourth aspect of the present invention has the
unexpected advantage of using common and cheaply available starting materials based
on 1-amino-8-hydroxynaphthylene disulphonic acid and thereby increases the breadth of
diazo components which may be used in the coupling reactions, thus significantly
increasing the synthetic versatility of the standard coupling process. In a further optional
25 step, the process comprises the complexation of a compound of Formula (1) with a metal,
preferably a transition metal.

30 A fifth aspect of the invention provides a compound of Formula (1) and salts thereof,
as hereinbefore defined, with the proviso that at least one of the groups represented by A
carries a group selected from sulpho and carboxy groups (sulpho do not include sulphato).
Preferably both groups represented by A carry a group selected from sulpho and carboxy
groups, more preferably both groups represented by A carry a sulpho group. Preferences
35 for A and L in the compounds of the fifth aspect of the invention are as hereinbefore defined
in relation to the first aspect of the invention.

35 A further aspect of the present invention provides a paper, an overhead projector
slide or a textile material printed with a composition, a compound or by means of a process
according to the present invention.

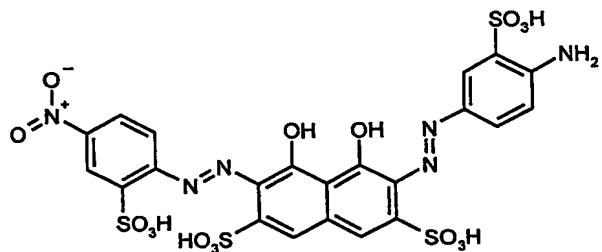
A still further aspect of the present invention provides an ink jet printer cartridge,
optionally refillable, comprising one or more chambers and a composition, wherein the
composition is present in at least one of the chambers, and the composition is as defined in
the second aspect of the present invention.

The present compounds and compositions provide prints of attractive, neutral black shades which are particularly well suited for the ink jet printing of text and images. The compositions have good storage stability and low tendency to block the very fine nozzles used in ink jet printers. Furthermore, the resultant images have good optical density, light-fastness, wet-fastness and resistance to fading in the presence of oxidising air pollutants (e.g. ozone).

5 The invention is illustrated by the following Examples in which all parts and percentages are by weight unless specified otherwise.

10 Example 1

Preparation of:



15 Stage (A) – Diazotisation and Coupling

5-Nitro-2-aminobenzene sulphonic acid (21.8 g, 0.10 moles) was added to water (300 ml), containing concentrated hydrochloric acid (30 g). Sodium nitrite (6.9 g, 0.10 moles) was added over 15 minutes at 0 to 5°C. After 60 minutes the excess nitrous acid was destroyed and to the resultant solution of diazonium salt was added slowly at 5 to 20 10°C and pH 1 to 2, a solution of 1-hydroxy-8-aminonaphthalene-3,6-disulphonic acid (31.9 g, 0.10 moles), which had been re-precipitated in water (500 ml).

25 2-Acetylamino-5-aminobenzene sulphonic acid (23.0 g, 0.11 moles) was added to water (300 ml), containing concentrated hydrochloric acid (32.6 g). Sodium nitrite (7.6 g) was added over 10 minutes at 0 to 5°C. After 60 minutes the resultant solution of diazonium salt was added over 120 minutes at 5 to 10°C and pH 8 to 9, to the above dye solution. After 10 hours the resultant product was precipitated by the addition of sodium chloride (100 g) and collected. Paste was then oven dried to give a dark blue solid (71g).

30 Stage (B) - Conversion of Amino to Hydroxy Group

The dark blue solid was dissolved in water (1000 ml) and heated to 80°C. Sodium hydroxide (40 g) was added and the temperature maintained at 70°C for 10 hours. After this time the pH was adjusted to pH 1 with concentrated hydrochloric acid and the solution allowed to cool to ambient temperature. The precipitate was then collected. The paste was then oven-dried to give 7.5 g of a black solid.

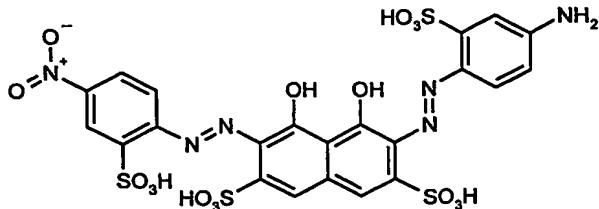
35 The black solid was dissolved in water (50 g) at pH 7 to 8 by the addition of sodium hydroxide (pearl). The solution was then dialysed using Visking™ tubing (<50

μScm^{-1}) and then screened through a cascade of filters (GF/A, GF/D, 0.45 μm) and dried in the oven to give the title product (7g) as a black solid.

Example 2

5

Preparation of:



5-Acetylaminobenzene sulphonic acid (23.0 g, 0.10 moles) was added to water (300 ml), containing concentrated hydrochloric acid (30 g). Sodium nitrite (6.9 g) was added over 10 minutes at 0 to 5°C. After 60 minutes the excess nitrous acid was destroyed and the resultant solution of diazonium salt was added slowly at 5 to 10°C and pH 8 to 9, to a solution of 1,8-dihydroxynaphthalene-3,6-disulphonic acid (32.0 g, 0.10 moles), which had been dissolved in water (500 g). The reaction was shown to have reacted quantitatively by HPLC.

15

5-Nitro-2-aminobenzene sulphonic acid (43.6 g, 0.20 moles) was added to water (500 g), containing concentrated hydrochloric acid (60 g). Sodium nitrite (13.8 g) was added over 15 minutes at 0 to 5°C. After 60 minutes the resultant solution of diazonium salt was added over 120 minutes at 5 to 10°C to the above dye solution to which previously had been added tetrahydrofuran (1000 g), whilst maintaining the pH at 7 by addition of solid Na_2CO_3 . After 5 hours the resultant precipitate was then collected and after oven drying gave a dark red solid (55.3 g).

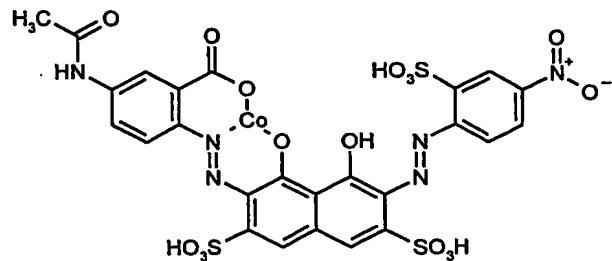
25

The above dark red solid was dissolved in water (1000 ml) and heated to 80°C. Sodium hydroxide (10 g) was added and the temperature maintained at 80°C for 8 hours. After this time the pH was adjusted to 7 to 8 with concentrated hydrochloric acid and the solution allowed to cool to ambient temperature. The solution was then dialysed using Visking™ tubing ($<50 \mu\text{Scm}^{-1}$) and then screened through a cascade of filters (GF/A, GF/D, 0.45 μm) and dried in the oven to give the title product as a black solid (47.2g).

Example 3

30

Preparation of:



5 A slurry of 5-acetylamino-2-aminobenzene carboxylic acid (64 g) and sodium nitrite (24 g) in water (500 ml) was added over 40 minutes to a mixture of hydrochloric acid (120 ml, 36°Tw) and ice (120 g). The resultant mixture was stirred at <5°C for 30 minutes and then excess nitrous acid was destroyed with sulfamic acid solution (10% w/v). The resulting solution was added over 1 hour to a solution of chromotropic acid (100 g) in water (800 ml) and ice (200 g) at pH 8 (sodium hydroxide, 48% w/v) at <5°C. During the addition the solution was maintained at pH 8-9.5. After a further hour the solution was allowed to warm to room temperature and stirred overnight.

10 The solution was adjusted to pH 1, filtered and the paste was washed with sodium chloride solution (500 ml, 17% w/v). The paste was then slurried in acetone (1000 ml) and dried in an oven to give a purple solid (152 g).

15 Cobalt acetate (11 g) was added in one go to a solution of the purple solid (35 g) in water (350 ml) at 80°C and pH 8.5 (concentrated ammonia solution) which was stirred for 2 hours. The solution was adjusted to pH 8.5 (concentrated ammonia solution) and allowed to cool to room temperature and stirred overnight.

20 Sodium chloride (35 g) was added to the solution which was stirred for one hour. A further portion of sodium chloride (35 g) was added and after one hour the mixture was filtered, washed with sodium chloride solution (300 ml, 20% w/v) and the resultant paste was slurried in acetone (500 ml) and collected by filtration. The paste was dried in an oven to give a dark blue solid (20 g).

25 A solution of 5-nitro-2-aminobenzene sulfonic acid (7 g) and sodium nitrite (2.5 g) in water (100 ml) at pH 8 (sodium hydroxide, 48% w/v) was added to a mixture of hydrochloric acid (20 ml, 36°Tw) and ice (20 g) over 20 minutes at <5°C. The mixture was stirred for 30 minutes and the excess nitrous acid was destroyed with sulfamic acid solution (10% w/v). This mixture was added dropwise over 35 minutes to a solution of the dark blue solid (18 g) in water (180 ml) and ice (20 g) at pH 8 (sodium hydroxide solution, 48% w/v) at <5°C. During the addition the solution was maintained at pH 8-9.5. The solution was stirred for a further hour at <5°C and then allowed to warm to room temperature overnight at pH 8.5.

30 Sodium chloride (45 g) was added and the solution was stirred for 2 hours. The resulting mixture was filtered and the collected paste was washed with sodium chloride solution (200 ml, 15% w/v). The paste was then slurried in acetone (500 ml), collected by filtration and slurried in acetone (300 ml) and collected. The resulting solid was dissolved in water (150 ml), dialysed using Visking™ tubing (<50 μ Scm⁻¹) and the resulting solution was screened through a cascade of filters (GF/A, GF/D, 0.45 μ m) and dried in the oven to give the title product as a black solid (9 g).

Examples 4 to 8

The method of Example 1 was repeated except that in place of 5-nitro-2-aminobenzene sulphonic acid there was used the amine listed in Table 1, column A and in place of 2-acetylaminobenzene sulphonic acid there was used the amine indicated in column B. For examples 4, 5 and 7, 1-hydroxy-8-aminonaphthalene-3,5-disulphonic acid was used in place of 1-hydroxy-8-aminonaphthalene-3,6-disulphonic acid. The resultant dyes had the formula shown in column C.

Table 1

10

Ex.	A	B	C	λ_{Max}
4				
5*				
6				
7*				
8 [#]				580nm

* synthesis included an additional hydrolysis step in aqueous sodium hydroxide to remove the acetyl group.

synthesis included an additional reduction step using sodium sulphite to convert the nitro group into an amino group.

15

Examples 9 to 44

The method of Example 2 was repeated except that in place of 5-acetylaminobenzene there was used the amine listed in Table 2, column A and in place of 5-nitro-2-aminobenzene sulphonic acid there was used the amine indicated in column B. 5

The resultant dye had the formula shown in column C. In examples 36, 37 and 38 the resultant dye is a metal complex, complexation was carried out, here as a final step, in an analogous manner to Example 3 above using copper(II) acetate, cobalt(II) acetate and nickel acetate respectively.

10

Table 2

Ex.	A	B	C	λ Max
9				556nm
10				
11				572nm
12*				
13				

14				
15				
16*				
17#				
18*				574nm
19				

20				576nm
21				
22*				
23				570nm
24				543nm
25				574nm

26*				568nm
27*				
28*				553nm
29				552nm
30*				573nm
31				570nm

32	<chem>OCCOc1ccc(cc1)Nc2ccc(cc2)O</chem>	<chem>NS(=O)(=O)c1ccc(cc1)N</chem>	<chem>OCCOc1ccc(cc1)N=Nc2ccc(cc2)Oc3ccc(cc3)N=Nc4ccc(cc4)S(=O)(=O)O</chem>	591nm
33	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)Oc2ccc(cc2)N=Nc3ccc(cc3)N=C1=NC(=O)O</chem>	
34	<chem>CC(=O)N1=NN=C1O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>CC(=O)N1=NN=C1Oc2ccc(cc2)N=Nc3ccc(cc3)S(=O)(=O)O</chem>	
35*	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>Nc1ccc(cc1)S(=O)(=O)Oc2ccc(cc2)N=Nc3ccc(cc3)N=C1=NC(=O)O</chem>	
36	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)Oc2ccc(cc2)N=Nc3ccc(cc3)O[Cu]([O-])=N4=NC(=O)C(=O)N4=NC(=O)C(=O)O</chem>	
37	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)Oc2ccc(cc2)N=Nc3ccc(cc3)O[Cu]([O-])=N4=NC(=O)C(=O)N4=NC(=O)C(=O)O</chem>	
38	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)O</chem>	<chem>NC1=NN=C1C(=O)O</chem>	<chem>CC(=O)Nc1ccc(cc1)S(=O)(=O)Oc2ccc(cc2)N=Nc3ccc(cc3)O[Ni]([O-])=N4=NC(=O)C(=O)N4=NC(=O)C(=O)O</chem>	

39				
40*				
41				
42*				
43				
44*				

* synthesis included an additional hydrolysis step in aqueous sodium hydroxide to remove the acetyl group.

5 # synthesis included an additional reduction step using sodium sulphite to convert the nitro group into an amino group.

Examples 45 to 48

The method of Example 3 was repeated except that in place of 5-acetylaminobenzene carboxylic acid there was used the amine listed in Table 3, column A and in place of 5-nitro-2-aminobenzene sulfonic acid there was used the amine indicated in column B, and in place of cobalt acetate there was used the acetate of the metal indicated. The resultant dye had the formula shown in column C.

Table 3

Ex.	A	B	C	λ_{Max}
45				541nm
46				
47				599nm
48				

10

Example 49 - Ink formulations

Inks may be prepared according to the formulation, wherein Dye is each of the dyes from the Examples 1 to 48 above:

15

2-Pyrrolidone	5 parts
Thiodiglycol	5 parts
Surfynol™ 465	1 part (from Air Products Inc., USA)
Dye	3 parts
Water	86 parts

Example 50 – Ink Jet Printing

The inks prepared in Example 49 may be incorporated into an empty cartridge of a Hewlett Packard DeskJet 550C™ ink jet printer and printed onto Xerox X Acid™ paper to give prints of excellent shade, light and ozone fastness.

Claims

5 1. A process for printing an image on a substrate comprising applying thereto a composition comprising a liquid medium and a compound of Formula (1):



wherein:

10 each A independently is optionally substituted aryl or heteroaryl; and L is an optionally substituted, optionally metallised 1,8-dihydroxynaphthylene group;

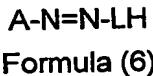
provided that at most one of the groups represented by A has a hydroxy substituent ortho to the -N=N- groups shown in Formula (1).

15 2. A composition comprising:

(a) 0.1 to 15 parts of the compound of Formula (1) as defined in claim 1; and
(b) from 85 to 99.9 parts of a liquid medium;

wherein all parts are by weight and the number of parts of (a)+(b)=100.

20 3. A process for preparing a compound of Formula (1), as defined in claim 1, which comprises diazotising an amine of formula A-NH₂ to give a diazonium salt, and coupling the resultant diazonium salt with a compound of Formula (6):



25 wherein L and each A independently are as defined in claim 1.

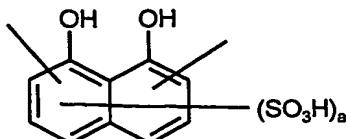
30 4. A process for the preparation of a compound of Formula (1) as defined in claim 1 which comprises the reaction of a compound of formula A-N=N-Q-N=N-A with a strong base, wherein each A independently is as defined in claim 1 and Q is an optionally substituted 1-hydroxy-8-amino-naphthylene group.

35 5. A compound of Formula (1), as defined in claim 1, with the proviso that at least one of the groups represented by A carries a group selected from sulpho and carboxy groups.

6. A compound of Formula (1) as defined in claim 1 wherein both groups represented by A carry a group selected from sulpho and carboxy.

40 7. A compound of Formula (1) as defined in claim 1 wherein both groups represented by A carry a sulpho group.

8. A compound of Formula (1) as defined in any of claims 5 to 7 wherein L is of Formula (2)



5 Formula (2)

wherein a is 1 or 2 and SO₃H is in free acid or salt form.

9. A paper, an overhead projector slide or a textile material printed by means of a process as defined in claim 1, with a composition as defined in claim 2, or with a compound 10 as defined in claim 5.

10. An ink jet printer cartridge, optionally refillable, comprising one or more chambers and a composition, wherein the composition is present in at least one of the chambers and the composition is as defined in claim 2.

THIS PAGE BLANK (USPTO)

